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(54) **Antigen binding proteins and methods for their production.**

(57) The present invention relates to a method for producing a protein corresponding to an antibody capable of binding to an antigen and to a protein prepared by this method, which protein is composed of predetermined framework regions of the heavy chain and light chain of an antibody, said predetermined regions being linked to undetermined regions which correspond in length to hypervariable regions of said antibody and which undetermined regions contain a sequence of amino acids capable of binding to said antigen. The present invention provides also various tools used in the said method. Furthermore, the present invention provides an antigen screening kit comprising a plurality of synthetic genes which may be used for screening antigens for binding to the proteins encoded by said synthetic genes.

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The present invention relates to a method for producing antigen binding proteins, which proteins are prepared by screening a library of synthetic genes containing randomized sequences after the expression of said genes with said antigen.

The very selective and specific binding characteristics of antibodies makes these molecules extremely attractive for use in a variety of medical and basic research applications. Traditional methods for generating antibodies involve immunization and hybridoma technology for the generation of monoclonal antibodies. Recently, polymerase chain reaction (PCR) based techniques have made it possible to engineer humanized antibodies which may serve as better therapeutic agents than their murine counterparts (Winter, and Milstein, 1991; Co, and Queen, 1991; Orlandi et al.1989). Furthermore, this technology has progressed to the point where it is now possible to clone the immunoglobulin (antibody) repertoire of an immunized mouse from spleen cells into phage expression vectors and identify expressed antibody fragments specific to the antigen used for immunization (Winter, and Milstein, 1991; Gussow et al.1989; Hodgson, 1991; Marks et al.1991; Garrard et al.1991; Duschosal et al.1992; Kang et al.1991b; Clackson et al.1991; Huse et al.1989; Persson et al.1991; Kang et al.1991a; Hoogenboom et al.1991; Barbas III et al. 1991). However, this technology has had little success in identifying specific antigen binding antibody fragments from unimmunized animals suggesting that there may be a prerequisite for prior immunization to the antigen of interest.

The present invention provides a method for producing a protein which binds to an antigen of choice, by using the antigen to screen a library of proteins which have been generated by using DNA synthesis and recombinant techniques combined with randomizing methods. The proteins, also referred to as synthetic antibodies, have the structure of antibodies, specifically Fab or Fv fragments, and contain randomized binding sequences which may correspond in length to hypervariable regions, i.e. complementary-determining regions (CDRs).

The techniques of this invention provide a method to generate a library of completely de novo synthesized antibody fragments which allows the bypass of both immunization and the necessity to use animals. The development of a synthetic antibody library has many advantages over other antibody libraries which are derived from immunized or unimmunized animals. The synthetic antibodies are developed without the use of animals (or hybridoma technology) and the problems associated with tolerance can be avoided. In addition, the synthetic antibody approach can be used for identifying antibodies against molecules which appear to be non-immunogenic or fail to induce immune response in animals. Furthermore, synthetic antibodies can be used to fill possible "holes" which may be present in an animals immune system repertoire.

The structure of an immunoglobulin consists of heavy and light chains which can be further defined into variable and constant domains which are indicated above. The smallest antibody fragment which forms an antigen binding site is referred to as an Fv fragment. Genetic engineering techniques have made it possible to generate single chain antibody (Fv) fragments. These Fv fragments consist of the heavy and light chain variable regions tethered together by a flexible glycine-serine linker. The variable regions can be further subdivided into framework regions which are fairly conserved among antibodies and hypervariable regions (CDR) which are quite diverse and are important in defining antigen specificity.

There are many uses for such synthetic antibodies and libraries. Some exemplary uses are listed below.

Synthetic antibody libraries can be used to complement other types of antibody libraries derived from animals in any drug screening or other ligand screening procedures.

Synthetic antibody libraries can be manipulated and modified for use in combinatorial type approaches in which the heavy and light chain variable regions are shuffled and exchanged between synthetic antibodies in order to affect specificities and affinities. This enables the production of antibodies which bind to a selected antigen with a selected affinity. For example, catalytic antibodies (abzymes) could be constructed. Antibodies with enhanced affinities can also be produced.

The approach of constructing synthetic single chain antibodies is directly applicable to constructing synthetic Fab fragments which can also be easily displayed and screened in the same manner.

The diversity of the synthetic antibody libraries can be increased by altering the chain lengths of the CDRs and also by incorporating changes in the framework regions which may affect antibody affinity. In addition, alternative libraries can be generated with varying degrees of randomness or diversity by limiting the amount of degeneracy at certain positions within the CDRs. The synthetic library can be modified further by varying the chain lengths of the CDRs and adjusting amino acids at defined position in the CDRs or the framework region which may affect affinities. Antibodies identified from the synthetic antibody library can easily be manipulated to adjust their affinity and or effector functions. In addition, the synthetic antibody library is amenable to use in combinatorial type approaches used by others. This may result in the

increasing of the affinities of the synthetic antibodies during the screening procedures.

The synthetic antibody library can be used for the generation and identification of anti-idiotypic antibodies which may mimic ligand and/or receptor molecules, and CDRs from screened synthetic antibodies can be used as potential peptidomimetics.

5 Screening of the synthetic antibody library can be modified to identify synthetic antibodies which may interact with their ligand under certain defined conditions (i.e., under reducing conditions which may be present in the intracellular environment.)

The strategy of constructing de novo synthetic antibodies can be adapted to the development of peptide libraries which are conformational in nature.

10 Synthetic antibodies identified from screening can be used for diagnostics such as the identification of any disease marker. Also, synthetic antibodies identified from screening can be used for the development of immunotherapeutics such as antibodies which can be administered for passive immunization or immunoconjugates which may be used to target tumors or other targets.

15 The coding sequences for identified synthetic antibodies can be manipulated using state of the art cloning strategies so that their antigen binding specificity can be grafted onto any immunoglobulin class or subtype.

The synthetic antibody library can be used for the screening of minute amounts of antigen which may not be available in sufficient quantity for the immunization of an animal.

20 The synthetic antibodies can be expressed to high levels in both prokaryotes and eukaryotes using present available technologies.

The synthetic antibodies can be used in any and all applications in which antibodies derived from other sources or by other means are used.

#### Brief Description of the Figures

- 25 Figure 1. A. Structure of a complete antibody molecule  
B. Structure of single-chain antibody molecule (Fv)  
Figure 2. Amino acid sequence of a synthetic Fv. Hypervariable region residues are replaced with X to represent any of the 20 amino acids.  
 30 Figure 3. Nucleotide sequence encoding a synthetic Fv as it is depicted in Fig. 2. n represents any nucleotide. Codon usage is biased for expression in E. coli and S. cerevisiae.  
Figure 4. Examples for the oligonucleotides synthesized for use in the generation of synthetic gene templates.  
Figure 5. Diagram of the PCR based production of synthetic genes encoding Fv.  
 35 Figure 6. Ethidium bromide stained agarose gel showing synthetic gene product of the second PCR step.  
Figure 7. Diagram of the FUSE 5 phage display vector.  
Figure 8. Diagram of Gene III phagemid vector BLSKDSg III.  
Figure 9. Diagram of the helper phage E. coli strains PJD1 and PJD2.  
 40 Figure 10. Diagram of fusion proteins displayed by phagemid and helper phage.  
Figure 11. Diagram of microorganisms displaying Fv antibodies.  
Figure 12. Antibody screening protocol, wherein panel A represents the incubation of phage/bacteria expressing synthetic Fv fragments with immobilized antigen. Panel B represents the washing of unbound and non-specific phage/bacteria from antigen. Panel C represents the elution of bound phage/bacteria from the antigen and the enrichment of the phage/bacteria through sequential rounds of screening.  
 45 Figure 13. Amino acid sequence of anti-tat Fv compared with sequence of Fig. 2.  
Figure 14. Nucleotide sequence of anti-tat Fv compared with sequence of Fig. 3.

This invention is directed to a method for producing a protein corresponding to an antibody capable of binding to an antigen as outlined in the appended claims. This method involves synthesizing a plurality of synthetic genes, each of which contains both a predetermined nucleotide region encoding the framework regions of portions of the heavy chain and light chain of an antibody and undetermined nucleotide regions which contain a random sequence of nucleotides. The proteins encoded by the synthetic genes are expressed by inserting vectors containing the synthetic genes into microorganisms and allowing expression to occur. The expressed proteins are screened by using the antigen to obtain the protein which is capable of binding to the antigen. In one variant of this method, an undetermined nucleotide region may correspond 55 of binding to the antigen. In one variant of this method, an undetermined nucleotide region may correspond in length to a nucleotide sequence which encodes a hypervariable region of an antibody to which the protein may correspond.

Synthetic genes, which are double-stranded oligonucleotides, may be assembled by any conventional method. DNA synthesis or recombinant techniques, or polymerase chain reaction or any combination of such techniques are contemplated.

A synthetic gene may be synthesized by providing plurality of oligonucleotides, each of which contains a portion of the synthetic gene. All the oligonucleotides when combined together form the entire nucleotide sequence of the synthetic gene, e.g., the predetermined and undetermined regions. The sequence of the oligonucleotides in combination is considered to include the sequence of either strand of the synthetic gene, which is double-stranded. For example, both the sequence of the coding strand and the sequence complementary thereto are included. The oligonucleotides themselves are synthesized by stepwise addition of nucleotides with the undetermined nucleotide regions that contain a random sequence of nucleotides being synthesized by stepwise addition of one nucleotide out of a mixture of nucleotides. A mixture of oligonucleotides may contain any two or more of the nucleotide bases adenine, guanine, cytosine, and thymine. Also included may be modified bases such as inosine. The mixture may be an equal mixture of any 2 or more bases or the mixture may contain predetermined fractions of any 2 or more bases, or the mixture may be completely random. The bases may be synthesized by known methods and are also commercially available from various suppliers of biochemical reagents. Synthesis as described above may be accomplished by attachment of bases to a solid substrate and sequential addition of an individual base from a vessel containing such base, or of an unknown base from a vessel containing the mixture described. This may be done by machine as described in the Example below. The synthetic gene is then synthesized by annealing and extending the plurality of oligonucleotides. Polymerase chain reaction (PCR) is one method for producing synthetic genes (see Example below). Any other method for assembling the synthesized oligonucleotides and to create either strand of the synthetic gene may be used.

In a preferred approach, the oligonucleotides are used as PCR primers to obtain a single-stranded template for the synthetic gene. Each oligonucleotide used contains portions of the predetermined and undetermined regions of the synthetic gene, as described above. In addition, each oligonucleotide contains at its 5' end and its 3' end a nucleotide sequence of about 20 bases which sequence is complementary to about 20 bases of the sequence adjoining the given oligonucleotide's sequence on the synthetic gene. Under conditions well-known to be suitable for PCR, the set of oligonucleotides will anneal and extend to form a final product which is a single-stranded sequence forming one strand of the synthetic gene. This template can be used to form the synthetic gene by any conventional means. The complementary strand can be produced by adding a primer, bases, and a polymerase, for example. For much more efficient production, PCR can be used. Primers corresponding to either end of the synthetic gene can be artificially synthesized by any conventional means (most of the sequence of the synthetic gene is already known, as described above, and therefore the primer sequences are easily deduced). These primers are added to the synthetic gene template which was obtained as described above, under PCR conditions, which are well-known in the art. The final product of this reaction are multiple copies of the synthetic gene. This full approach as described may be used to form a plurality of synthetic genes, each gene containing a different undetermined region with a different specificity.

The vectors and microorganisms used to express the synthetic genes may be any conventional vectors and microorganisms. Examples are provided infra.

This invention also is drawn to a plurality of proteins, each protein being composed of predetermined framework regions of portions of the heavy and light chain of an antibody, which are linked to undetermined regions of the antibody, and which contain a random sequence of amino acids. The length of these undetermined regions may be any desired length. A preferred length is a length corresponding to that of a hypervariable region of an antibody. At least one of the proteins is capable of binding to an antigen for which an antibody is sought. The proteins may be single chain proteins or may be composed of more than one polypeptide chain. A specific example is a single chain protein capable of binding to HIV-1 tat protein. Said single-chain protein is composed of predetermined framework regions of portions of the heavy-chain and light-chain of an antibody, which are linked to undetermined regions, which regions correspond in length to hypervariable regions of said antibody. The undetermined regions contain a sequence of amino acids capable of binding to HIV-1 tat protein. A preferred embodiment of this protein has the amino acid sequence

5      Glu Val Gln Leu Val Glu Ser Gly Arg Gly Leu Val Gln Pro Gly Gly  
       Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Phe  
       Leu Val Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
       Ala Thr Tyr Ser Met Ile Ser Arg Ala Arg Val Leu Asp Gly Ser Phe  
       Asn Gly Arg Tyr Thr Ile Ser Arg Asp Asp Ser Lys Asn Thr Leu Tyr  
       Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
       Ala Arg Ile Gly Ser Thr His Thr Ile Pro Arg Leu Ser Gln Tyr Gly  
       Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Ser Gly  
 10      Gly Gly Gly Ser Gly Gly Gly Gly Ser Asp Ile Gln Met Thr Gln Ser  
       Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys  
       Lys Leu Arg Gly Pro Gln Pro His Ala Ile Thr Trp Tyr Gln Gln Lys  
       Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Tyr Asp Gly Gln Thr Leu  
       Val Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe  
       Thr Pro Thr Ile Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr  
 15      Cys Thr Pro Thr His Lys Ile Asp Ser Pro Phe Gly Gln Gly Thr Lys  
       Val Glu Ile Lys Arg Thr      [SEQ ID NO:11]

shown in Figure 13.

20      A further aspect of the present invention is a synthetic gene which encodes a single-chain polypeptide capable of binding to HIV-1 tat protein as described above. A particular example is the synthetic gene comprising the nucleotide sequence of Figure 14.

25      GAAGTTCAAC TGGTTGAATC CGGTCGTGGT CTGGTTCAAC CAGGTGGTTC CCTGCGTCTG  
       TCCTGTGCTG CTTCCGGTTT CACCTTCTCC CATTTTTTGG TGGCGTGGGT TCGTCAAGCT  
       CCAGGTAAAG GTCTGGAATG GGTTGCTACC TACTCAATGA TTAGCCGGGC CCGAGTACTC  
 30      GATGGCTCCT TTAATGGACG TTACACCATC TCCCGTGACG ACTCCAAAAA CACCCTGTAC  
       CTGCAAATGA ACTCCCTGCG TGCTGAAGAC ACCGCTGTTT ACTACTGTGC TCGTATTGGT  
       TCTACGCACA CAATCCCACG ACTGTCTCAA TACGGGGGTC AAGGTACCCT GGTTACCGTT  
 35      TCTCCGGTG GTGGTGGTTC CGGTGGTGGT GGTTCTGGTG GTGGTGGTTC CGACATCCAA  
       ATGACCCAAT CCCCATCCTC TCTGTCCGCT TCCGTTGGTG ACCGTGTTAC CATCACCTGT  
       AAACCTCAGAG GACCACAACC ACACGCCATT ACATGGTACC AACAAAAACC AGGTAAAGCT  
       CCAAACTGC TGATCTACTA CGACGGCCAA ACGTTGGTGG GTGTTCCATC CCGTTTCTCC  
 45      GGTTCTGGTT CTGGTACCGA CTTACCCCG ACCATCTCCT CTCTGGAACC AGAAGACTTC  
       GCTACCTACT ACTGTACTCC TACGCACAAG ATCGATAGCC CATTCGGTCA AGGTACCAAA  
       GTTGAAATCA AACGTACC      [SEQ ID NO:12]  
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55      The predetermined nucleotide regions of the synthetic gene encode selected regions of an antibody. Both the heavy and light chain subunits of an antibody are made up of conserved regions and variable regions, as is well known in the art. The variable regions themselves contain framework regions which themselves are relatively conserved, and complementary-determining (CDR) or hypervariable regions which are not conserved and which are specific to a given antibody. These regions determine binding specificity. The synthetic genes are designed to encode framework regions from both heavy and light chain variable regions, interspersed with undetermined regions containing random amino acid sequences. The undeter-

mined regions may be of any length, and length may be selected to give desired effects. The length of the undetermined regions may correspond to the length of hypervariable regions of an antibody, such that the undetermined regions "fill in" for hypervariable regions and provide a randomized selection of possible binding specifications and affinities. The framework regions are derived from known antibodies. The boundaries of framework and hypervariable regions are well known in the art and one skilled in the art can determine the regions by conventional means. It is possible to obtain such antibodies from hybridomas, a variety of which are available commercially from depositories such as the American Type Culture Collection (ATCC), 12301 Parklawn Drive, Rockville, Maryland U.S.A. or from biological supply houses. Hybridomas can also be produced by conventional methods. Or antibodies may be obtained from any cells which naturally express them or have gene inserts enabling their expression. Genes encoding antibodies may be obtained from any such sources and from cells which contain but do not express antibody genes. Actual antibodies or antibody genes may be used to make the synthetic genes with well known techniques of protein synthesis or genetic engineering. However, a preferred alternative is to obtain the known sequences of numerous specific antibodies from scientific publications, from patent publications or from a computer database such as those provided by Genbank or Brookhaven National Labs. A consensus framework sequence can then be generated based on these sequences. An example of such a sequence is the amino acid sequence

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20  Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
    Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Xaa Xaa
    Xaa Xaa Xaa Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Trp Val
    Ala Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
    Xaa Xaa Arg Phe Thr Ile Ser Arg Asp Asp Ser Lys Asn Thr Leu Tyr
25  Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
    Ala Arg Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
    Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly
    Gly Gly Gly Ser Gly Gly Gly Gly Ser Asp Ile Gln Met Thr Gln Ser
    Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys
    Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Trp Tyr Gln Gln Lys
30  Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Xaa Xaa Xaa Xaa Xaa Xaa
    Xaa Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Thr Asp Phe
    Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr
    Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Phe Gly Gln Gly Thr Lys
35  Val Glu Ile Lys Arg Thr [SEQ ID NO:1],

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the sub-parts of which sequence are specified in Figure 2. In this sequence the amino acids of the hypervariable regions have been replaced with "X" to represent any amino acid. Such a sequence may be synthesized by conventional methods as described above. As indicated above such a sequence obviates the need to use antibodies of animal origin and the limitations of such use, and the need to use cell cultures and cloning to obtain antibodies. Further, the consensus sequence can be biased to favor expression in a selected microorganism, as described in the Examples. The synthetic genes may encode any antibodies or parts thereof. Preferred synthetic genes encode proteins which correspond to antibodies known in the art as Fv antibodies. These antibodies are composed of the variable regions (including hypervariable and framework) of the light and heavy chains of an antibody which are connected to each other by peptide bonds via a peptide linker sequence to form a single-chain polypeptide. Also preferred are synthetic genes encoding the antibodies known in the art as Fab fragments. These antibodies are also composed of the heavy and light chain variable regions, but form a double chain polypeptide wherein the heavy and light chain segments may be connected by disulfide bridges.

Part of this invention is a plurality of synthetic genes which encodes the plurality of proteins, each of which synthetic genes contains nucleotide sequences which encode predetermined framework regions of portions of the heavy chain and light chain of an antibody, which are linked to nucleotide sequences which encode undetermined regions containing a random sequence of amino acids. These regions may contain a number of nucleotides which encode a sequence corresponding in length to hypervariable regions of an antibody. At least one protein of the plurality of proteins is capable of binding to an antigen for which an antibody is desired. A particular example is a plurality of proteins wherein each of the proteins is a single-chain protein, or is composed of more than one chain.

Also contemplated in the invention is a vector having inserted therein a synthetic gene which contains nucleotide sequences encoding predetermined framework regions of portions of the heavy chain and light chain of an antibody, linked to nucleotide sequences which encode undetermined regions of any desired length, which also may correspond in length to hypervariable regions of an antibody, and containing a random sequence of amino acids. This vector is capable of causing expression of the synthetic gene as a protein by a microorganism. A preferred vector is one which can cause the expressed protein to translocate to the outer surface of the microorganism which contains the synthetic gene. Vectors include any conventional vectors. Vectors such as plasmids, cosmids, viruses, transposons, and any other elements capable of genetic transfer are contemplated. The synthetic genes are inserted into the vectors by methods well known in the art of genetic engineering. Vectors capable of causing expression are intended to include all conventional genetic elements for inducing gene expression, e.g., start and stop codons, promoters, enhancers, etc. Vectors which can cause expression of a protein on the surface of a microorganism may include signal sequences which cause the protein to go through a cell membrane. Microorganism can be any cells into which vectors may be inserted. However, phages are herein also considered to be microorganisms. Display phages can be used into which genes may be inserted in such a position as to be expressed as a fusion protein with one of the phage's coat proteins on the phage's surface. Microorganisms may be bacteria, such as *E. coli*, yeast, fungi, algae, mammalian cells or any other prokaryotic or eukaryotic cell whether acellular or part of a tissue.

All vectors and microorganisms described are conventional and well-known in the art. Also conventional are techniques for inserting synthetic genes into vectors, and for inserting vectors into microorganisms. Transformation, transfection, electroporation, and protoplast fusion are examples of well-known methods.

Also part of this invention is a plurality of microorganisms, each of which has on its outer surface at least one protein of the plurality of proteins, each of the proteins being composed of predetermined framework regions of portions of the heavy-chain and light-chain of an antibody, which are linked to undetermined regions of any length, in particular corresponding length to hypervariable regions of the antibody and containing a random sequence of amino acids. Any conventional microorganism, such as a phage, may be used. Vectors such as those described above may be used to insert the synthetic genes which encode the proteins. The phages may themselves include such a synthetic gene. At least one of these proteins is capable of binding to an antigen for which an antibody is desired to be found. The plurality of microorganisms may be used as a screen to determine which of the proteins expressed by the microorganism that binds to a predetermined antigen. Any conventional screening method may be used. For example, the antigen may be fixed to a solid support such as a culture dish or a bead in a column. Medium containing the plurality of proteins expressed the surfaces of microorganisms is contacted with the support. The protein capable of binding to the antigen will bind to the immobilized antigen itself and thereby will itself be immobilized. Then, unbound protein is washed off. Next, the bound protein, still attached to the microorganism expressing it, is eluted from the antigen. Washing and elution conditions are well known in the art. The isolated microorganism contains the synthetic gene which encodes the protein which binds to the antigen. This synthetic gene can be used in conventional recombinant technology to produce the antigen binding protein in quantity and also in any desired modified forms. For example, the synthetic gene can be expressed in company with genes expressing constant regions of an antibody, under conditions known to cause aggregation of the protein with the constant regions to produce a complete antibody. Heavy chain constant regions of IgM, IgG, IgA, IgD, or IgE types could be used. Light chain constant regions of kappa or lambda types would also be used to combine with the protein.

Alternatively, the proteins themselves may be isolated from the microorganisms and used for screening by conventional means.

Furthermore the present invention provides an antigen screening kit comprising a plurality of synthetic genes which may be used for screening antigens for binding to the protein encoded by said synthetic genes.

The synthetic genes, the vectors comprising them as well as microorganisms transformed with said synthetic genes or vectors may be used in a diagnostic test system for detecting various parameters which may be useful for a physician.

Moreover the proteins prepared in accordance with the present invention may be used as a diagnostic test in place or in combination with regular antibodies such as monoclonal or polyclonal antibodies.

The Example which follows further describes the invention but is not intended to limit the invention in any way.

Example

PCR is used to generate a library of completely de novo synthetic single chain antibodies (Fv) which consist of the heavy and light chain variable regions tethered together by a flexible glycine-serine linker (Fig. 1). Using a compilation of known human immunoglobulin amino acid sequences, a synthetic single chain Fv antibody fragment which contains conserved framework residues found in human antibodies and random residues in the hypervariable regions was designed. These artificial variable heavy and light chain domains are joined by a glycine-serine linker which for correct folding of the synthetic Fv fragment to allow formation of antigen binding sites. The synthetic Fv amino acid sequence was then reverse translated into a nucleic acid sequence with codon usage biased for expression in *E. coli*. The amino acids of the hypervariable regions were represented by degenerate triplets (NNN). The DNA encoding the synthetic Fv molecules was generated by a modification of the gene construction PCR method (Dillon, and Rosen, 1990). The resultant de novo synthesized Fv PCR products have been cloned into phage and phagemid display vectors or into bacterial outer membrane protein fusion expression vectors. In the phage display vector the single chain Fv is expressed as a fusion protein with the coat III protein of a M13 derivative single strand DNA bacteriophage (FUSE 5) (Parmley, and Smith, 1988). In the PAL (peptidoglycan associated lipoprotein) fusion vector, the Fv fragment should be expressed on the outer surface of the *E. coli* outer membrane as a fusion within the PAL protein (Fuchs et al.1991; Chen, and Henning, 1987). Expression of the Fv fragments in phage or bacteria should allow for the rapid screening of the library by incubation of expressing phage or bacteria with immobilized antigen and sequential enrichment of specific antigen binding Fv expressing phage or bacteria. Since the DNA encoding the synthetic Fv will be present in the enriched phage or bacteria it is possible to sequence and subclone the single chain Fv fragments into additional antibody expression vectors.

These synthetic antibody libraries are screened with various antigens which have been immobilized on coated dishes, magnetic beads and affi-gel columns. The successful development and screening of these libraries allows the generation of novel antibody fragments, without the use of animals (or hybridoma technology), which recognize a wide variety of molecules including, non-immunogenic and tolerant epitopes, transcription factors, nuclear components, lipids, carbohydrates, etc. By virtue of the random amino acid sequence built into the hypervariable regions, the synthetic Fv library has the potential to bind almost any antigen regardless of its immunogenicity.

**Design of synthetic single chain antibody sequence:** A compilation of known human antibody sequences was used to generate a consensus amino acid sequence of the variable regions for the light chain based on Kabat subgroup I and the heavy chain based on Kabat subgroup III (Kabat et al.1987). Residues contained within the hypervariable regions (CDRs) for the heavy and light chains were replaced with X amino acid, where X can represent any of the twenty amino acids. The redesigned heavy and light variable region sequences were then bridged by a flexible linker sequence encoded by the sequence

Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser  
[SEQ ID NO:13].

The resulting synthetic antibody amino acid sequence [SEQ ID No: 1] shown in Figure 2 was then reverse translated into the nucleic acid sequence



GAAGTTCAAC TGGTTGAATC CGGTGGTGGT CTGGTTCAAC CAGGTGGTTC CCTGCGTCTG  
 TCCTGTGCTG CTTCCGGTTT CACCTTCTCC NNNNNNNNNN NNNNNTGGGT TCGTCAAGCT  
 5 CCAGGTAAAG GTCTGGAATG GGTGCTNNN NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN  
 NNNNNNNNNN NNNNNNNNCG TTTCACCATC TCCCGTGACG ACTCCAAAAA CACCCTGTAC  
 10 CTGCAAATGA ACTCCCTGCG TGCTGAAGAC ACCGCTGTTT ACTACTGTGC TCGTNNNNNN  
 NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN NNNTGGGGTC AAGGTACCCT GGTACCGTT  
 TCCTCCGGTG GTGGTGGTTC CGGTGGTGGT GGTTCGGTG GTGGTGGTTC CGACATCCAA  
 15 ATGACCCAAT CCCCATCCTC TCTGTCCGCT TCCGTTGGTG ACCGTGTAC CATCACCTGT  
 NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN NNNTGGTACC AACAAAAACC AGGTAAAGCT  
 CCAAACTGC TGATCTACNN NNNNNNNNNN NNNNNNNNNG GTGTTCCATC CCGTTTCTCC  
 20 GGTTCGGTT CTGGTACCGA CTTACCCTG ACCATCTCCT CTCTGCAACC AGAAGACTTC  
 GCTACCTACT ACTGTNNNNN NNNNNNNNNN NNNNNNNNNN NNTTCGGTCA AGGTACCAA  
 25 GTTGAATCA AACGTACC [SEQ ID NO:2]

shown in Figure 3. In this nucleotide sequence the codon usage is biased for expression in *E. coli* and for expression in *S. cerevisiae*. The degenerate X amino acid residues were encoded using degenerate codons of nnn where n represents any of the four nucleotides A,C,G or T.

30 **PCR technique for generating DNA encoding synthetic single chain antibody sequence:** DNA encoding the synthetic antibody sequence was generated using an adaptation and modification of the method described by Dillon and Rosen (Dillon, and Rosen, 1990) for the PCR construction of synthetic genes and is outlined in Figure 5. Briefly, eight long oligonucleotides having the following nucleotide sequences

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GAAGTTCAAC TGGTTGAATC CGGTGGTGGT CTGGTTCAAC CAGGTGGTTC CTGCGTCTG  
 TCCTGTGCTG CTTCCGGTTT CACCTTCTCC NNNNNNNNNN NNNNNTGGGT TCGTCAAGCT  
 5 CCAGG [SEQ ID NO:3]

GGAGTCGTCA CGGGAGATGG TGAAACGNNN NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN  
 10 NNNNNNNNNN NNNNNNNNAG CAACCCATTC CAGACCTTTA CCTGGAGCTT GACGAACCCA  
 [SEQ ID NO:4]

CGTTTCACCA TCTCCCGTGA CGACTCCAAA AACACCCTGT ACCTGCAAAT GAACTCCCTG  
 15 CGTGCTGAAG ACACCGCTGT TTACTACTGT GCTCGT [SEQ ID NO:5]

CACCGGAGGA AACGGTAACC AGGGTACCTT GACCCCANNN NNNNNNNNNN NNNNNNNNNN  
 20 NNNNNNNNNN NNNNNNACGA GCACAGTAGT AACAGCGGT G [SEQ ID NO:6]

TGGTTACCGT TTCCTCCGGT GGTGGTGGTT CCGGTGGTGG TGGTTCTGGT GGTGGTGGTT  
 25 CCGACATCCA AATGACCCAA TCCCCATCCT CTCTGTCCGC TTCCGTTGGT GACCGTGTTA  
 CCATCA [SEQ ID NO:7]

GATCAGCAGT TTTGGAGCTT TACCTGGTTT TTGTTGGTAC CANNNNNNNN NNNNNNNNNN  
 30 NNNNNNNNNN NNNNNNACAGG TGATGGTAAC ACGGTCACCA ACGGAA [SEQ ID NO:8]

GGTACGTTTG ATTTCAACTT TGGTACCTTG ACCGAANNNN NNNNNNNNNN NNNNNNNNNN  
 35 NNNACAGTAG TAGGTAGCGA AGTCTTCTGG TTGCAGAGAG GAGATGGTCA GGGTGAAGT  
 40 [SEQ ID NO:9]

CAGGTAAAGC TCCAAACTG CTGATCTACN NNNNNNNNNN NNNNNNNNNN GGTGTTCCAT  
 45 CCCGTTTCTC CGGTTCCGGT TCTGGTACCG ACTTCACCCT GACCATCTCC TCTCTG  
 [SEQ ID NO:10]

were synthesized on an ABI oligonucleotide synthesizer which spanned the designed sequence of the synthetic antibody. These oligonucleotides were between 100 and 135 nucleotides in length and contained

short overlaps approximately 20 nucleotides in length (Fig. 4). The overlaps were positioned such that they corresponded to defined sequences of the framework regions. Nucleotide positions designated by *n* were synthesized such that any of the four (A,C,G,T) phosphoramidites would be introduced to the solid support at the same time. This was accomplished by having the four phosphoramidites premixed in solution and placed in a separate reservoir which was utilized during synthesis for base positions designated *n*. Flanking primers were also synthesized which contained appropriate restriction sites to facilitate cloning.

Briefly, the two step PCR approach was used for generating the DNA fragment. The first PCR step was used to generate the full length templates and conditions were as follows 0.5 micrograms of each of the eight long overlapping oligonucleotides were mixed in a 100 microliter PCR reaction containing 2.5 units of AmpliTaq DNA polymerase and subjected to 35 cycles of thermal cycling in a Perkin-Elmer 9600 System thermal cycler. Cycle conditions were as follows: 5 minute initial denaturation at 94 °C ; 15 seconds at 94 °C , 15 seconds at 55 °C , 45 seconds at 72 °C for 35 cycles, followed by a final extension at 72 °C for 3 minutes. A second PCR reaction was used to generate material for cloning. One to three microliters of the product from the first PCR reaction was used as template for a second reaction containing one microgram of each flanking primer and subjected to 25 cycles of thermal cycling as described above.

#### Vectors:

The phage display vector, FUSE 5 (Fig. 7) was used for cloning the single chain antibody DNA in frame with the amino terminus of the gene III phage coat protein DNA at engineered SfiI sites (Parmley, and Smith, 1988).

The phagemid display vector BLSKDSgenelII (Fig. 8) was constructed by ligating the lac promoter from the pDS56 vector (Bujard et al.1987) as an XhoI-SphI fragment and a synthetic pelB leader sequence as a SphI-PstI fragment into the XhoI-PstI sites of Bluescript SK+ (Stratagene, La Jolla, California). The resultant plasmid, BLSKDSpelB was further manipulated to include genelII as an XbaI-NotI fragment which was obtained by PCR cloning from M13mp18 (New England Biolabs, Beverly, Massachusetts).

The peptidoglycan associated lipoprotein (PAL) bacterial display vector BLSKDSPAL was constructed by PCR cloning of the PAL sequence (Chen, and Henning, 1987) from *E. coli* strain MC1061 (BioRad, Richmond, California) using a 5' primer which contained BamHI, NsiI, and XbaI sites and a 3' primer which contained a NotI site. The PAL PCR product was then cloned as a BamHI-NotI fragment into BLSKDSpelB.

#### Construction of *E. coli* helper phage strains:

**PJD1 (Fig. 9):** MC1061 was cotransformed with the lac repressor expression vector pDM1.1 (Bujard et al.1987) and single strand DNA from the FUSE 2 phage (a tetracycline transducing phage obtained from George Smith) (Parmley, and Smith, 1988). The PJD1 strain is tetracycline and kanamycin resistant and can be made transformation competent for both heat shock and electroporation.

**PJD2 (Fig. 9):** Similar to PJD1 but lacks the pDM1.1 plasmid.

**PJD3:** MC1061 transformed by the interference resistant helper phage VCSM 13 (Stratgene). This strain is kanamycin resistant.

**Construction of antibody phage libraries:** The synthetic single chain antibody PCR products were digested at their termini with SfiI and ligated into the SfiI sites of the FUSE5 phage display vector. Four micrograms of cut vector DNA was mixed with 0.5 micrograms of cut insert and ligated in a final volume of one milliliter with 5 units of T4 ligase and incubated at 16 °C for twelve hours. Ligations were then ethanol precipitated and resuspended in 10 microliters of water. The ligation mixture was then electroporated into electrocompetent MC1061 cells using a Biorad electroporator set at 2.5 kV, 400 ohms and 25 microfarads. The cells were then resuspended in 2 mL of SOC medium (20 g/l bacto-tryptone, 5 g/l bacto-yeast extract, 0.5 g/l NaCl, 2.5mM KCl, 10mM MgCl<sub>2</sub>, 20mM glucose; adjusted to pH 7) and incubated in Falcon 2071 polystyrene tube for one hour at 37 °C. The transformed cells were then plated on LB agar plates containing 25 microgram per mL tetracycline and incubated overnight at 37 °C. Tetracycline resistant colonies were then scraped from the plates into TBS (50mM Tris-HCl pH 7.5, 150mM NaCl). Phages expressing the antibody were isolated and concentrated by poly(ethylene glycol) (PEG) precipitation, which was performed as follows: Pellet phage culture at 4000 rpm for 15 minutes at 4 degrees C (Beckman JA10 rotor or equivalent). Pour supernatant into clean bottle and precipitate phage by adding PEG 8000 4% w/v and NaCl to 3% w/v to the supernatant. Shake for about 5 minutes to dissolve. Incubate on ice for 30 minutes. Pellet phage at 9000 rpm for 20 minutes at 4 degrees C (Beckman JA10 or equivalent). Resuspended phage pellets in TBS.

**Construction of antibody phagemid libraries:** The synthetic single chain antibody PCR products were

digested at their termini with NsiI and XbaI and ligated into the PstI and XbaI sites of the BLSKDSgenIII display vector. Ligation mixtures were then electroporated into either electrocompetent *E. coli* strains PJD1, PJD2 or PJD3 which contain helper phage (as described). Transformed cells were selected on 1.5% LB agar (10 g/l bacto-tryptone, 5 g/l bacto-yeast extract, 5 g/l NaCl, 15 g/l agar) containing ampicillin (100  $\mu$ g/mL), tetracycline (25  $\mu$ g/mL) and IPTG. Phage was then prepared by scraping colonies and treating as described above.

**Construction of antibody PAL libraries:** The synthetic single chain antibody PCR products were digested at their termini with NsiI and XbaI and ligated into the NsiI and XbaI sites of the BLSKDSPAL display vector. Ligation mixes were electroporated as described above and transformed bacteria was grown on 1.5% LB amp agar (10 g/l bacto-tryptone, 5 g/l bacto-yeast extract, 5 g/l NaCl, 15 g/l agar) plates overnight. Colonies were then scraped and stored as glycerol stocks at -70 °C until use in screening.

### Screening Protocols (Fig. 12)

Antigens used in screening were coupled to tosylactivated M-280 magnetic beads according to known methods. Dynabeads M-280 are uniform superparamagnetic polystyrene beads which may be obtained from Dynal, Great Neck, New York. Beads with similar properties from the same or other suppliers could also be used. Antigens were also immobilized on Nunc 96 well micotiter plates or affigel resin for use in screening antibody phage (phagemids, or PAL fusion bacteria).

Screening using coated magnetic beads was carried out in siliconized microfuge tubes which had been preincubated with TBS plus 1% BSA for one hour at room temperature. For primary screenings, one  $\mu$ L of antigen coated beads were mixed with 5  $\mu$ L of antibody phage preparations in a final volume of 1 mL of TBS plus 0.1% Tween-20 and 1% BSA. Incubations were carried out at 4 °C for one hour. The Phage bound magnetic beads were then concentrated using a MPC-6 (Dynal) magnetic particle concentrator and unbound phage was aspirated. The beads were then washed 10 to 20 times with TBS plus 0.1% Tween-20. This was done to wash away residual unbound and nonspecific phage. Phage which remained bound to the beads following the wash procedure were then eluted with either low pH, 0.2N HCl or by treatment with trypsin. In the case of low pH elution, the eluted phage were removed from the beads and neutralized with 2M Tris. The eluted phage were then used for infecting starved K91kan cells (a male *E. coli* strain obtained from George Smith) (Parmley, and Smith, 1988). The phage infected cells were selected for ampicillin (BLSKDS gene III phagemid Antibody library) or tetracycline (FUSE 5 Antibody library) transducing units. In the case of the phage library, antibody phage particles were prepared directly from the transduced colonies and used for sequential rounds of screening as described above. The phagemid library required the rescue procedure described next.

### Rescue of phagemid:

**Phagemid rescue procedures:** Antibody phagemid infected K91kan cells were scraped from plates and grown in liquid culture for one hour at 37 °C at which time the culture was divided in half. One half was used for preparing phagemid DNA by the alkaline lysis procedure while the other half was used for rescue by use of either FUSE2 or VCSM 13 helper phage. The rescue was achieved by adding  $10^8$  helper phage to the K91kan cells and incubating for an additional hour at 37 °C. After one hour the IPTG (final concentration = 1mM) was added and in the case of FUSE2, tetracycline was also added. The culture was incubated for 4 to 8 hours at which time the culture supernatant was used to prepare packaged phagemid for sequential rounds of screening.

An alternative approach for phagemid rescue used transformation of the PJD1, PJD2, and PJD3 strains by the isolated phagemid DNA. In this procedure, the transformed strains were selected with the appropriate antibiotics and rescued phagemid was prepared as described above and used in sequential screenings.

### Results

The initial phage library constructed contained approximately  $10^6$  to  $10^7$  independent clones. This library was screened against magnetic beads coated with the HIV-1 tat protein. The results of one screening is shown in Table I. One phage TR5 was identified which appeared to bind specifically to the Tat protein. In this experiment incubation of the purified phage with increasing amounts of tat coated beads resulted in an increase in the number of bound phage while little phage was observed to bind to increasing amounts of beads coated with other proteins.

**Table I**  
**ANTIBODY PHAGE TR5 SCREEN**

**First round screen**

Input Phage =  $10^{10}$  Phage Particles from FUSE5 Synthetic FV Library  
Screened with 1 microliter of HIV-1 Tat Protein Coated Magnetic Beads  
Five tet<sup>r</sup> Colonies were Obtained

**Second round screen**

Phage TR5 was Grown and Screened against Tat and other Protein Coated Beads

PROTEIN	TAT	GP120	p65
1	277	1	0
5	1639	4	ND
10	4800	14	ND

Magnetic Bead  
Volume  
(microliters)

Number of tet<sup>r</sup> colonies

In addition, the TR5 phage was selected for in 4 separate screening experiments as determined by DNA sequencing of the phage insert of antibody phage which enriched against Tat protein.

Sequence comparison of the phage TR5 insert and the initial framework sequence designed showed few differences between the two as indicated in figure 15. These changes did not significantly alter the amino acid sequence as compared in figure 14. It is unclear if these alterations are a result of PCR amplification of the initial construct or subsequent PCR cloning steps or if they arose as a result of mutation in the phage genome. Characterization of the unscreened library indicated a selective pressure or stability constraints against some insert sequences as evidenced by observations of partial and entire deletions of the antibody insert.

Results from the phagemid library indicate that the insert is more readily maintained and that the cloning efficiency for construction of the library is much higher thereby making it possible to generate a larger and more diverse library. Our experiments show that it is possible to generate packaged phagemid by direct transformation into the PJD helper E.coli strains. Initial screening has shown that the same phagemid has been picked up 4 times and each of the 4 phage contain the same partial sequence.

The construction of a library composed of entirely synthetic antibodies has the potential to generate antibody molecules which have completely novel binding characteristics and the ability to bind virtually any antigen available for screening. Known human antibody sequences are used to form a consensus type framework sequence on which to base the design of an exemplary single chain antibody sequence. Following the design of the amino acid sequence of the synthetic antibody, the amino acid sequence was reverse translated into a nucleic acid sequence which contained codons that should be preferentially utilized

in *E. coli*. This differs significantly from previously published antibody libraries which have all been derived from animal tissue (Garrard et al.1991; Kang et al.1991b; Kang et al.1991a; Persson et al.1991; Huse et al.1989; Barbas III et al.1991; Gussow et al.1989; Clackson et al.1991; McCafferty et al.1990; Marks et al.1991; Hoogenboom et al.1991; Winter, and Milstein, 1991; Hodgson). The representation and expression of specific antibodies from these libraries may be hindered due to little or no expression of some library members as a result of poor codon usage in *E. coli*. The problems of codon usage in *E. coli* may be important for the generation of good libraries since phage (or phagemid) based vectors are being used for the display of the antibody molecules. Therefore, the inherent characteristic of the DNA encoding the synthetic antibody (SYNAB jargon term) may lead to increased expression of our antibody library.

The identification of the phage TR5 and its ability to bind the Tat protein confirm that functional synthetic antibodies have been generated based on a comparative analysis of known antibody sequences. This approach may be applied to the study of other proteins which belong to larger families such as the T cell receptors.

The use of phage display vectors offers many options for the screening of large antibody libraries. Screening conditions can be altered to select for various affinities. The use of gene III as a fusion for the single chain antibody allows expression of a limited number of molecules which may lower nonspecificity during screening. The use of trypsin for elution results in increased recovery of phage compared to low pH. The use of trypsin does not seem to interfere with the infectivity of the phage.

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10 antibodies by repertoire cloning. *Proc. Natl. Acad. Sci. USA* 88, 2432-2436.
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## SEQUENCE LISTING

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 (I) TELEX: 962292/965542 hlr ch

(ii) TITLE OF INVENTION: Antigen binding proteins and  
 Methods for their production

(iii) NUMBER OF SEQUENCES: 13

## (iv) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk  
 (B) COMPUTER: IBM PC compatible  
 (C) OPERATING SYSTEM: PC-DOS/MS-DOS  
 (D) SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)

## (vi) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 07/843,125  
 (B) FILING DATE: 28-FEB-1992

## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 246 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

Glu	Val	Gln	Leu	Val	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	1	5	10	15
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Xaa	Xaa	20	25	30	
Xaa	Xaa	Xaa	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	35	40	45	
Ala	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	Xaa	50	55	60	



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5 Xaa Xaa Arg Phe Thr Ile Ser Arg Asp Asp Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Trp  
 100 105 110  
 Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly  
 115 120 125  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Asp Ile Gln Met Thr Gln Ser  
 130 135 140  
 Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys  
 145 150 155 160  
 Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Trp Tyr Gln Gln Lys  
 165 170 175  
 Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Xaa Xaa Xaa Xaa Xaa Xaa  
 180 185 190  
 Xaa Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe  
 195 200 205  
 Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr  
 210 215 220  
 Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Phe Gly Gln Gly Thr Lys  
 225 230 235 240  
 Val Glu Ile Lys Arg Thr  
 245

(2) INFORMATION FOR SEQ ID NO:2:

35 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 738 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

40 (ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

45 GAAGTTCAAC TGGTTGAATC CGGTGGTGGT CTGGTTCAAC CAGGTGGTTC CCTGCGTCTG 60  
 TCCTGTGCTG CTTCCGGTTT CACCTTCTCC NNNNNNNNNN NNNNNTGGGT TCGTCAAGCT 120

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TCAGGTAAAG GTCTGGAATG GGTGCTNNN NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN 180  
 NNNNNNNNNN NNNNNNNNCG TTTCACCATC TCCCGTGACG ACTCCAAAAA CACCCTGTAC 240  
 CTGCAAATGA ACTCCCTGCG TGCTGAAGAC ACCGCTGTTT ACTACTGTGC TCGTNNNNNNN 300  
 NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN NNNTGGGGTC AAGGTACCCT GGTACCGTT 360  
 TCCTCCGGTG GTGGTGGTTC CGGTGGTGGT GGTCTGGTG GTGGTGGTTC CGACATCCAA 420  
 ATGACCCAAT CCCCATCCTC TCTGTCCGCT TCCGTTGGTG ACCGTGTTAC CATCACCTGT 480  
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 CCAAACTGC TGATCTACNN NNNNNNNNNN NNNNNNNNNG GTGTTCCATC CCGTTTCTCC 600  
 GGTCCGGTT CTGGTACCGA CTTACCCCTG ACCATCTCCT CTCTGCAACC AGAAGACTTC 660  
 GCTACCTACT ACTGTNNNNN NNNNNNNNNN NNNNNNNNNN NNTTCGGTCA AGGTACCAA 720  
 GTTGAATCA AACGTACC 738

(2) INFORMATION FOR SEQ ID NO:3:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 125 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GAAGTTCAAC TGGTTGAATC CGGTGGTGGT CTGGTTCAAC CAGGTGGTTC CCTGCGTCTG 60  
 TCCTGTGCTG CTTCCGGTTT CACCTTCTCC NNNNNNNNNN NNNNNTGGGT TCGTCAAGCT 120  
 CCAGG 125

(2) INFORMATION FOR SEQ ID NO:4:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 120 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

GGAGTCGTCA CGGGAGATGG TGAAACGNNN NNNNNNNNNN NNNNNNNNNN NNNNNNNNNN 60

5 NNNNNNNNNN NNNNNNNNAG CAACCCATTC CAGACCTTTA CCTGGAGCTT GACGAACCCA 120

## (2) INFORMATION FOR SEQ ID NO:5:

10 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 96 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

15 (ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

20 CGTTTCACCA TCTCCCGTGA CGACTCCAAA AACACCCTGT ACCTGCAAAT GAACTCCCTG 60

CGTGCTGAAG ACACCGCTGT TTACTACTGT GCTCGT 96

## (2) INFORMATION FOR SEQ ID NO:6:

25 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 101 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

30 (ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

35 CACCGGAGGA AACGGTAACC AGGGTACCTT GACCCCANNN NNNNNNNNNN NNNNNNNNNN 60

NNNNNNNNNN NNNNNNACGA GCACAGTAGT AAACAGCGGT G 101

## (2) INFORMATION FOR SEQ ID NO:7:

40 (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 126 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

45 (ii) MOLECULE TYPE: cDNA

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## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

TGGTTACCGT TTCCTCCGGT GGTGGTGGTT CCGGTGGTGG TGGTTCTGGT GGTGGTGGTT 60  
 5 CCGACATCCA AATGACCCAA TCCCCATCCT CTCTGTCCGC TTCCGTTGGT GACCGTGTTA 120  
 CCATCA 126

## (2) INFORMATION FOR SEQ ID NO:8:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 106 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

GATCAGCAGT TTTGGAGCTT TACCTGGTTT TTGTTGGTAC CANNNNNNNNN NNNNNNNNNN 60  
 NNNNNNNNNN NNNNNACAGG TGATGGTAAC ACGGTCACCA ACGGAA 106

## (2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 119 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

GGTACGTTTG ATTTCAACTT TGGTACCTTG ACCGAANNNN NNNNNNNNNN NNNNNNNNNN 60  
 NNNACAGTAG TAGGTAGCGA AGTCTTCTGG TTGCAGAGAG GAGATGGTCA GGGTGAAGT 119

## (2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 116 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

CAGGTAAAGC TCCAAACTG CTGATCTACN NNNNNNNNNN NNNNNNNNNN GGTGTTCCAT 60  
 5 CCCGTTTCTC CGGTTCCGGT TCTGGTACCG ACTTCACCCT GACCATCTCC TCTCTG 116

## (2) INFORMATION FOR SEQ ID NO:11:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 246 amino acids  
 (B) TYPE: amino acid  
 (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

Glu Val Gln Leu Val Glu Ser Gly Arg Gly Leu Val Gln Pro Gly Gly  
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 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Phe  
 20 20 25 30  
 Leu Val Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ala Thr Tyr Ser Met Ile Ser Arg Ala Arg Val Leu Asp Gly Ser Phe  
 50 55 60  
 Asn Gly Arg Tyr Thr Ile Ser Arg Asp Asp Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Ile Gly Ser Thr His Thr Ile Pro Arg Leu Ser Gln Tyr Gly  
 100 105 110  
 Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly  
 115 120 125  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Asp Ile Gln Met Thr Gln Ser  
 130 135 140  
 Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys  
 145 150 155 160  
 Lys Leu Arg Gly Pro Gln Pro His Ala Ile Thr Trp Tyr Gln Gln Lys  
 165 170 175  
 Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Tyr Asp Gly Gln Thr Leu  
 180 185 190

EP 0 557 897 A1

Val Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe  
 195 200 205  
 Thr Pro Thr Ile Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr  
 210 215 220  
 Cys Thr Pro Thr His Lys Ile Asp Ser Pro Phe Gly Gln Gly Thr Lys  
 225 230 235 240  
 Val Glu Ile Lys Arg Thr  
 245

(2) INFORMATION FOR SEQ ID NO:12:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 738 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: cDNA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

GAAGTTCAAC TGGTTGAATC CGGTCGTGGT CTGGTTCAAC CAGGTGGTTC CCTGCGTCTG 60  
 TCCTGTGCTG CTTCCGGTTT CACCTTCTCC CATTTTTTGG TGGCGTGGGT TCGTCAAGCT 120  
 CCAGGTAAAG GTCTGGAATG GGTGCTACC TACTCAATGA TTAGCCGGGC CCGAGTACTC 180  
 GATGGCTCCT TTAATGGACG TTACACCATC TCCCGTGACG ACTCCAAAAA CACCCTGTAC 240  
 CTGCAAATGA ACTCCCTGCG TGCTGAAGAC ACCGCTGTTT ACTACTGTGC TCGTATTGGT 300  
 TCTACGCACA CAATCCCACG ACTGTCTCAA TACGGGGGTC AAGGTACCCT GGTTACCGTT 360  
 TCCTCCGGTG GTGGTGGTTC CGGTGGTGGT GGTTCTGGTG GTGGTGGTTC CGACATCCAA 420  
 ATGACCCAAT CCCCATCCTC TCTGTCCGCT TCCGTTGGTG ACCGTGTTAC CATCACCTGT 480  
 AAACTCAGAG GACCACAACC ACACGCCATT ACATGGTACC AACAAAAACC AGGTAAAGCT 540  
 CCAAACTGC TGATCTACTA CGACGGCCAA AC GTTGGTGG GTGTTCCATC CCGTTTCTCC 600  
 GGTTCTGGTT CTGGTACCGA CTTACCCCG ACCATCTCCT CTCTGGAACC AGAAGACTTC 660  
 GCTACCTACT ACTGTACTCC TACGCACAAG ATCGATAGCC CATTCCGTCA AGGTACCAAA 720  
 GTTGAAATCA AACGTACC 738

## (2) INFORMATION FOR SEQ ID NO:13:

## (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 15 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: peptide

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser  
 1 5 10 15

## Claims

1. A method for producing a protein corresponding to an antibody capable of binding to an antigen which method comprises:
  - a.) synthesizing a plurality of synthetic genes, each of said synthetic genes containing a predetermined nucleotide region encoding the framework regions of portions of the heavy chain and light chain of said antibody, and undetermined nucleotide regions which contain a random sequence of nucleotides;
  - b.) causing the expression of the plurality of proteins encoded by all of the synthetic genes by microorganisms having inserted therein vectors containing said synthetic genes; and
  - c.) screening said plurality of expressed proteins to obtain said protein capable of binding to the antigen.
2. The method of claim 1 wherein an undetermined nucleotide region corresponds in length to a nucleotide sequence which encodes a hypervariable region of the antibody to which the protein corresponds.
3. The method of claim 1 wherein the synthetic gene included within said plurality of synthetic genes is synthesized by providing a plurality of oligonucleotides each of which contains a portion of the nucleotide sequence of a synthetic gene, the plurality of oligonucleotides being constructed such that all of said oligonucleotides combined together form the entire undetermined and determined nucleotide region sequence of said synthetic gene or a sequence complementary thereto, said oligonucleotides being synthesized by the stepwise addition of nucleotides, with the undetermined nucleotide regions which contain a random sequence of nucleotides being synthesized by the stepwise addition of one nucleotide from a mixture of nucleotides, and said synthetic gene being synthesized by annealing and extending said plurality of oligonucleotides to form said synthetic gene.
4. The method of claim 3 wherein the predetermined nucleotide regions of said oligonucleotides are synthesized stepwise by adding one of the individual nucleotides adenine, cytosine, guanine, or thymine and the undetermined nucleotide regions of said oligonucleotides are synthesized stepwise by addition of any one of said nucleotides from a mixture.
5. The method of claim 3 wherein the plurality of oligonucleotides are annealed and extended by a polymerase chain reaction.
6. The method of claim 4 wherein the undetermined nucleotide regions correspond in length to a nucleotide region which encodes the hypervariable regions of the antibody.
7. The method of claim 1 wherein the vectors containing said synthetic genes are display vectors.

8. The method of claim 6 wherein said plurality of proteins expressed by the microorganisms are located at the surface of said microorganisms through the use of said display vector, with said screening of said plurality of proteins for binding to said antigen, being carried out while said plurality of proteins are located at the surface of said microorganisms.
- 5 9. A plurality of proteins, each of said proteins being composed of predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being linked to undetermined regions which correspond in length to hypervariable regions of said antibody and which undetermined regions contain a random sequence of amino acids, at least one of said proteins being  
10 capable of binding to an antigen.
10. The plurality of proteins of claim 9 wherein each of said proteins is a single-chain protein.
11. The plurality of proteins of claim 9 wherein each of said proteins is composed of more than one  
15 polypeptide chain.
12. A single-chain protein capable of binding to an antigen and being composed of predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being linked to undetermined regions which correspond in length to hypervariable regions of  
20 said antibody and which undetermined regions contain a sequence of amino acids capable of binding to said antigen.
13. A protein as claimed in claim 12 capable of binding to HIV-1 tat protein and being composed of predetermined framework regions of portions of the heavy chain and light chain of an antibody, said  
25 predetermined regions being linked to undetermined regions which correspond in length to hypervariable regions of said antibody and which undetermined regions contain a sequence of amino acids capable of binding to HIV-1 tat protein.
14. A protein as claimed in claim 13 comprising the amino acid sequence [SEQ ID No: 11] shown in Figure  
30 13.
15. A synthetic gene which encodes a single-chain protein as claimed in claim 12 capable of binding to an antigen, which synthetic gene contains nucleotide sequences which encode predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being  
35 linked to nucleotide sequences which encode undetermined regions which correspond in length to hypervariable regions of said antibody, which undetermined regions contain a sequence of amino acids capable of binding to said antigen.
16. A synthetic gene which encodes a single-chain protein as claimed in claim 13 capable of binding to  
40 HIV-1 tat protein, which synthetic gene contains nucleotide sequences which encode predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being linked to nucleotide sequences which encode undetermined regions which correspond in length to hypervariable regions of said antibody, which undetermined regions contain a sequence of amino acids capable of binding to HIV-1 tat protein.
- 45 17. A synthetic gene as claimed in claim 16 comprising the nucleotide sequence [SEQ ID No: 12] shown in Figure 14.
18. A plurality of synthetic genes which encodes a plurality of proteins as claimed in any one of claims 9 to  
50 11, each of which synthetic genes contain nucleotide sequences which encode predetermined framework regions of portions of the heavy chain and light chain of an antibody linked to nucleotide sequences which encode undetermined regions which correspond in length to hypervariable regions of an antibody and which undetermined regions contain a random sequence of amino acids, at least one protein of said plurality of proteins being capable of binding to an antigen.
- 55 19. A vector capable of causing expression of a protein according to any one of claims 12 to 14 by a microorganism, said vector having inserted therein a synthetic gene which contains nucleotide sequences which encode predetermined framework regions of portions of the heavy chain and light chain



of an antibody linked to nucleotide sequences which encode undetermined regions which correspond in length to hypervariable regions of an antibody and which undetermined regions contain a random sequence of amino acids.

- 5 20. A vector of claim 19 wherein the vector which has the synthetic gene inserted therein is capable of causing expression of a protein in a microorganism and has the ability to cause translocation of a protein thus expressed to the outer surface of the microorganism.
21. A microorganism which contains a vector as claimed in claim 19 or 20.
- 10 22. An E. coli cell which contains a vector as claimed in claim 19 or 20.
23. A plurality of microorganisms each having inserted therein a vector capable of causing expression of a protein according to any one of claims 12 to 14 on the outer surface of the microorganism, each of said  
15 vectors containing a synthetic gene which encodes at least one protein of a plurality of proteins, each of said proteins being composed of predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being linked to undetermined regions which correspond in length to hypervariable regions of said antibody and which undetermined regions contain a random sequence of amino acids, at least one of said proteins being capable of binding to an  
20 antigen.
24. A phage or a phagemid which expresses on its outer surface a protein according to any one of claims 12 to 14 composed of predetermined framework regions of portions of the heavy chain and light chain of an antibody, said predetermined regions being linked to undetermined regions which correspond in  
25 length to hypervariable regions of said antibody and which undetermined regions contain a random sequence of amino acids.
25. An antigen screening kit comprising a plurality of synthetic genes as claimed in claim 18.
- 30 26. Use of a plurality of synthetic genes as claimed in claim 18 for screening antigen for binding to the proteins encoded by said synthetic genes.
27. The invention as hereinbefore described.

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Fig. 1/14

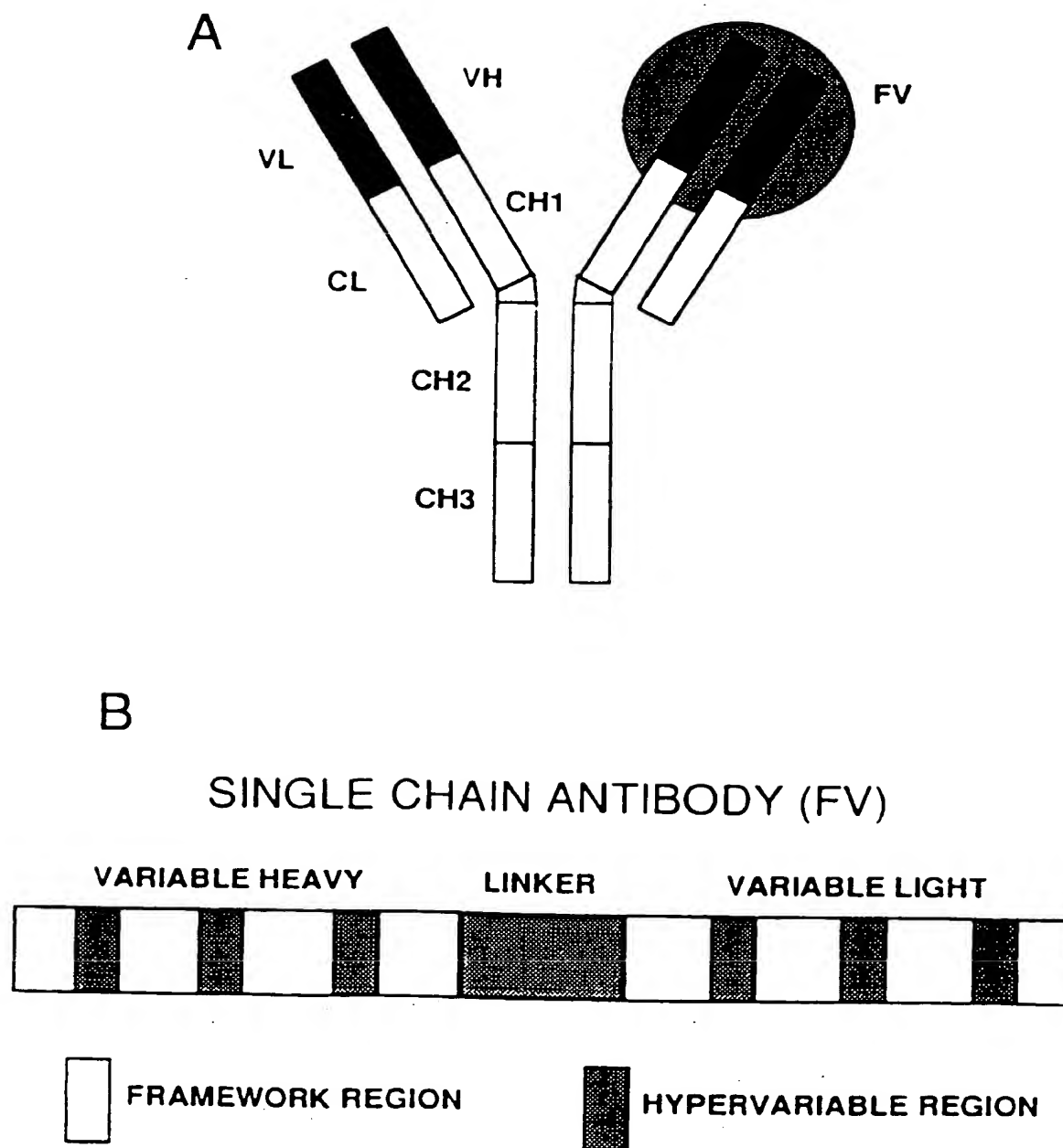


Fig. 2/14

AMINO ACID SEQUENCE OF SYNTHETIC FV

*Variable Heavy Region*

E V Q L V E S G G G L V Q P G G S L R L S  
 C A A S G F T F S [REDACTED] W V R Q A P G  
 K G L E W V A [REDACTED]  
 [REDACTED] R F T I S R D D S K N T L Y L Q M N  
 S L R A E D T A V Y Y C A R [REDACTED]  
 [REDACTED] W G Q G T L V T V S S

*Linker Region*

G G G G S G G G G S G G G G S

*Variable Light Region*

D I Q M T Q S P S S L S A S V G D R V T I  
 T C [REDACTED] W Y Q Q K P G K  
 A P K L L I Y [REDACTED] G V P S R F S  
 G S G S G T D F T L T I S S L Q P E D F A  
 T Y Y C [REDACTED] F G Q G T K V E  
 I K R T

Fig. 3/14

```

1  gaagttcaac  tgggtgaatc  cgggtggtggt  ctggttcaac  cagggtggttc
51  cctgcgtctg  tcctgtgctg  cttccggttt  caccttctcc  nnnnnnnnnn
101 nnnnntgggt  tcgtcaagct  ccaggtaaag  gtctggaatg  ggttgctnnn
151 nnnnnnnnnn  nnnnnnnnnn  nnnnnnnnnn  nnnnnnnnnn  nnnnnnnncg
201 tttcaccatc  tcccgtgacg  actccaaaaa  caccctgtac  ctgcaaataa
251 actccctgcg  tgctgaagac  accgctgttt  actactgtgc  tcgtnnnnnn
301 nnnnnnnnnn  nnnnnnnnnn  nnnnnnnnnn  nnntggggtc  aagggtaccct
351 gggtaccggt  tcctccggtg  gtggtggttc  cgggtggtggt  ggttctggtg
401 gtggtggttc  cgacatccaa  atgacccaat  ccccatcctc  tctgtccgct
451 tccgttggtg  accgtgttac  catcacctgt  nnnnnnnnnn  nnnnnnnnnn
501 nnnnnnnnnn  nnntgggtacc  aacaaaaacc  aggtaaagct  ccaaaactgc
551 tgatctacnn  nnnnnnnnnn  nnnnnnnnng  gtgttccatc  ccgtttctcc
601 gggtccggtt  ctggtaccga  cttcacctg  accatctcct  ctctgcaacc
651 agaagacttc  gctacctact  actgtnnnnn  nnnnnnnnnn  nnnnnnnnnn
701 nnttcggtca  aggtaccaaa  gttgaaatca  aacgtacc

```

## Fig. 4/14

1 gaagttcaac tgggtgaatc cgggtggtggt ctggttcaac caggtggttc  
 51 cctgcgtctg tctgtgctg ctcccggtt caccttctcc nnnnnnnnnn  
 101 nnnntgggt tcgtcaagct ccagg [SEQ ID NO:3]

1 ggagtcgtca cgggagatgg tgaaacgnnn nnnnnnnnnn nnnnnnnnnn  
 51 nnnnnnnnnn nnnnnnnnnn nnnnnnnnag caaccattc cagacctta  
 101 cctggagctt gacgaacca [SEQ ID NO:4]

1 cgtttcacca tctcccgta cgactccaaa aacaccctgt acctgcaaat  
 51 gaactccctg cgtgctgaag acaccgctgt ttactactgt gctcgt  
 [SEQ ID NO:5]

1 caccggagga aacggttaacc aggttacctt gacccannn nnnnnnnnnn  
 51 nnnnnnnnnn nnnnnnnnnn nnnnnnacga gcacagtagt aaacagcgg  
 101 g [SEQ ID NO:6]

1 tggttaccgt ttcctccggt ggtggtggtt ccggtggtgg tggttctggt  
 51 ggtggtggtt ccgacatcca atgacccaa tccccatct ctctgtccgc  
 101 ttcggtggt gaccgtgtta ccatca [SEQ ID NO:7]

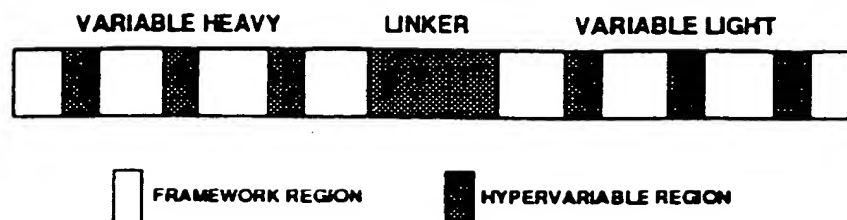
1 gatcagcagt tttggagctt tacctggtt ttgttggtac cannnnnnnn  
 51 nnnnnnnnnn nnnnnnnnnn nnnnnacagg tgatggtaac acggtcacca  
 101 acggaa [SEQ ID NO:8]

1 ggtacgtttg atttcaactt tggtagcttg accgaannnn nnnnnnnnnn  
 51 nnnnnnnnnn nnnacagtag taggtagcga agtcttctgg ttgcagagag  
 101 gagatggtca gggtagaagt [SEQ ID NO:9]

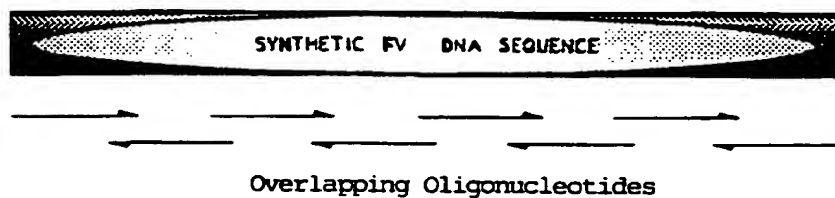
1 caggtaaagc tccaaaactg ctgatctacn nnnnnnnnnn nnnnnnnnnn  
 51 ggtgttccat cccgtttctc cggttccggt tctggtaccg acttcaccct  
 101 gaccatctcc tctctg [SEQ ID NO:10]

Fig. 5/14

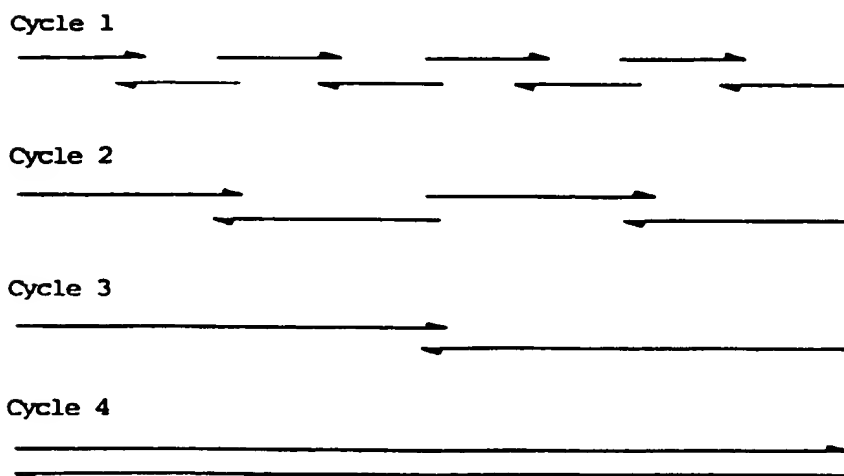
A. SINGLE CHAIN ANTIBODY (FV)



B. PRIMER DESIGN



C. FIRST PCR REACTION



D. SECOND PCR REACTION

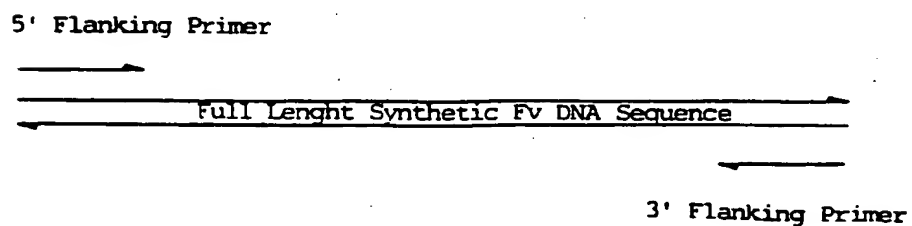


Fig. 6/14

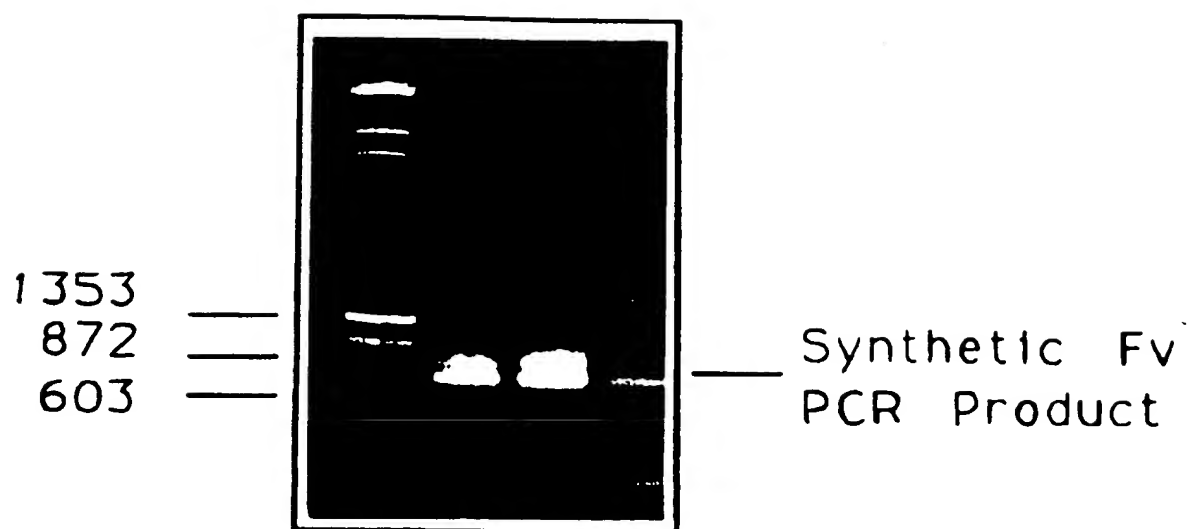


Fig. 7/14

**FUSE 5 PHAGE DISPLAY VECTOR**

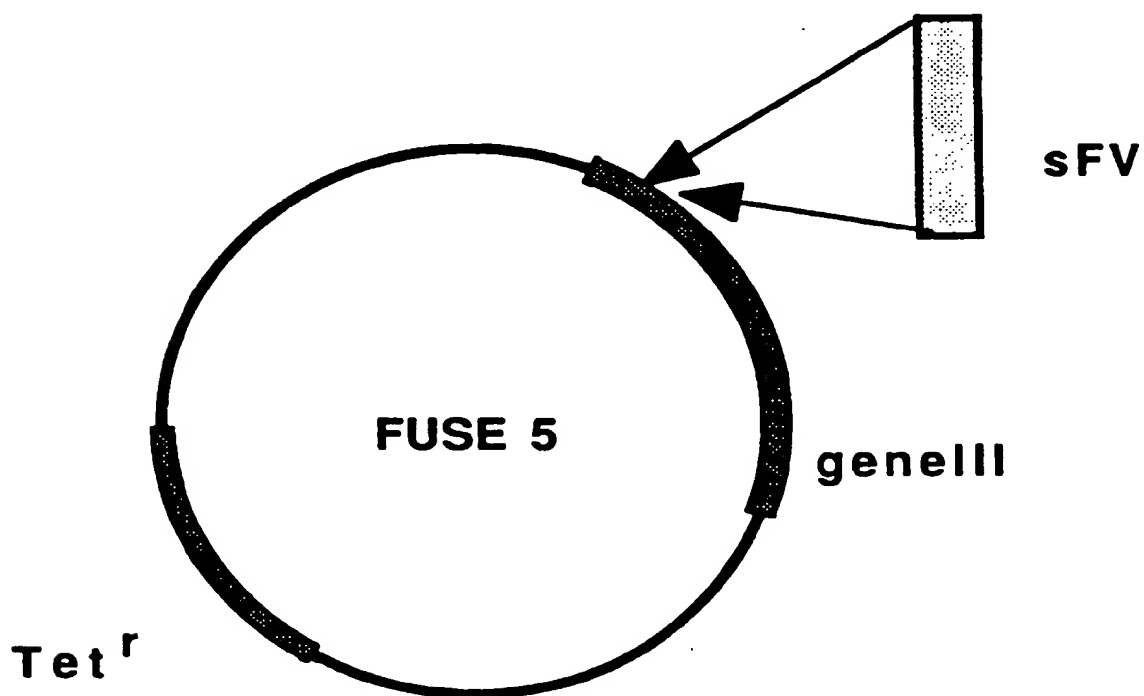




Fig. 8/14

**GENE III PHAGEMID VECTOR**

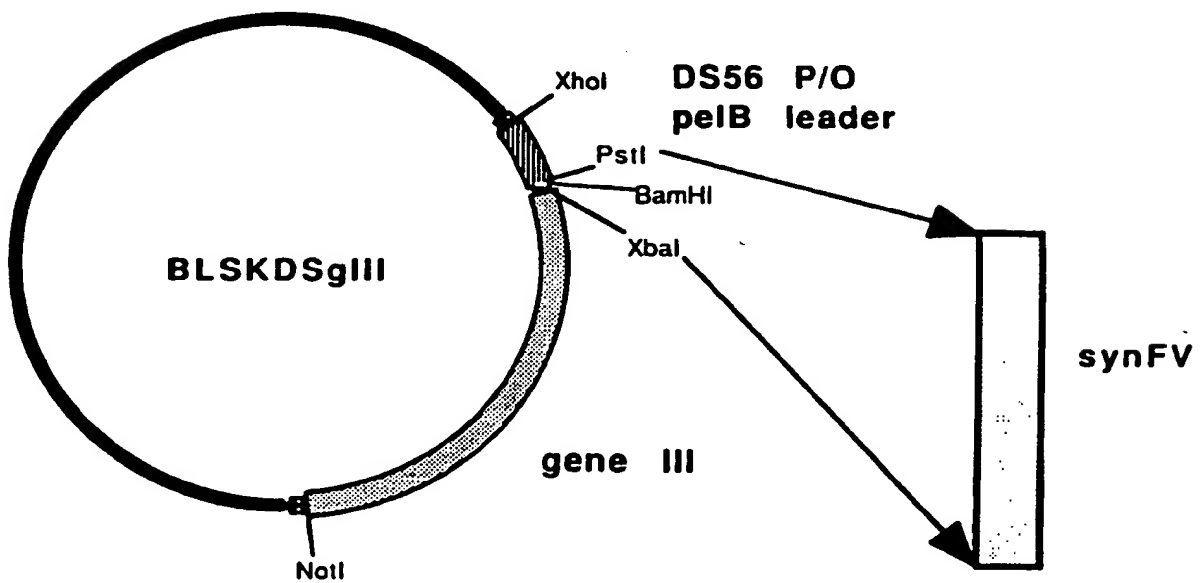


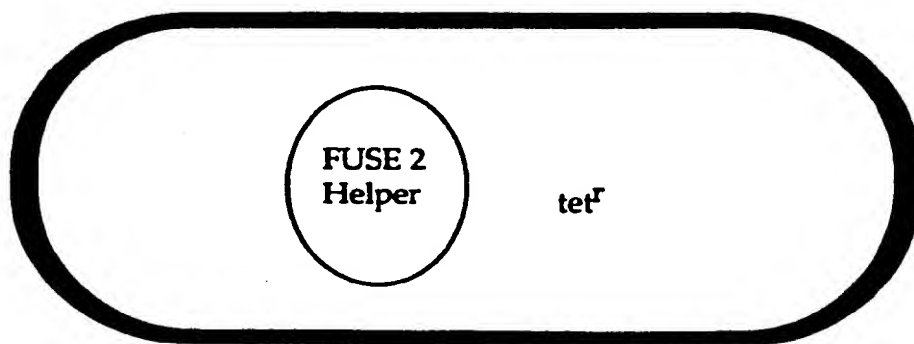
Fig. 9/14

**HELPER PHAGE E. coli STRAINS**



**PJD1 Strain**

MC1061+pDM1.1+FUSE 2



**PJD2 Strain**

MC1061+FUSE 2

Fig. 10/14

**Packaged Phagemid and Helper Phage Particles  
Will Display Fusion Proteins**

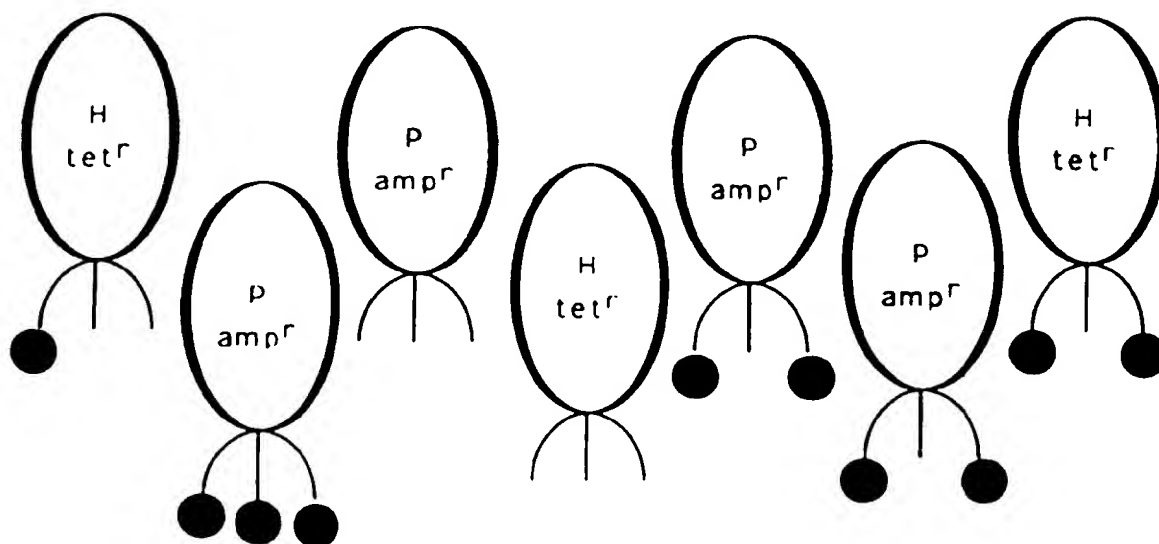
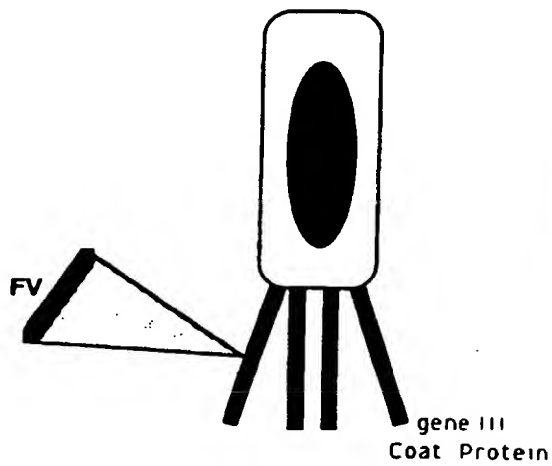
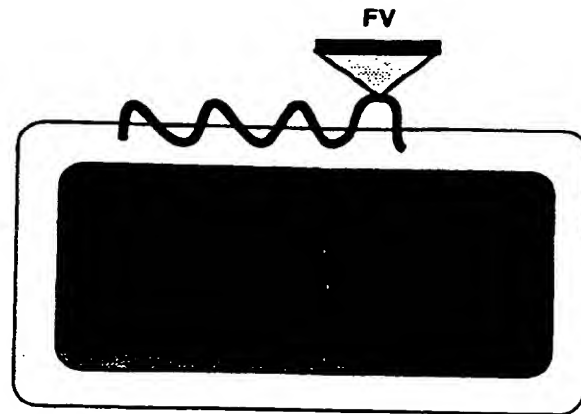


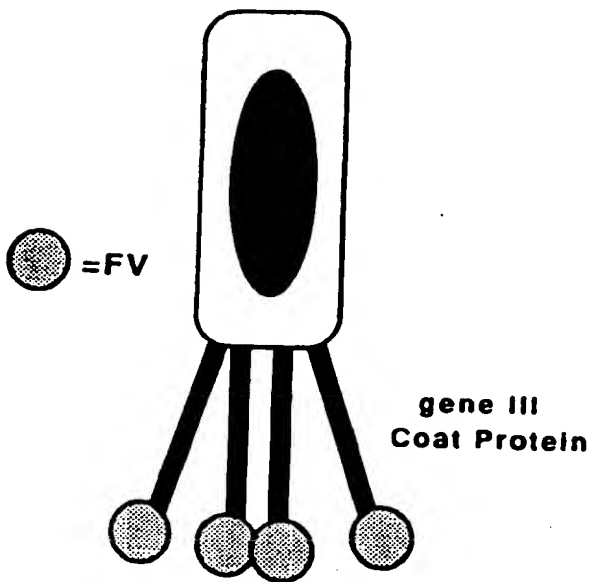
Fig. 11/14



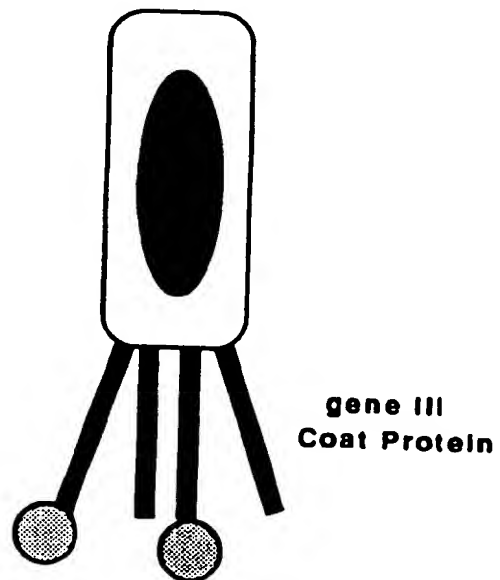
PHAGE ANTIBODY LIBRARY



BACTERIAL ANTIBODY FUSION LIBRARY



PHAGE ANTIBODY LIBRARY



PHAGEMID ANTIBODY LIBRARY

Fig. 12/14

**ANTIBODY LIBRARY SCREENING PROTOCOL**

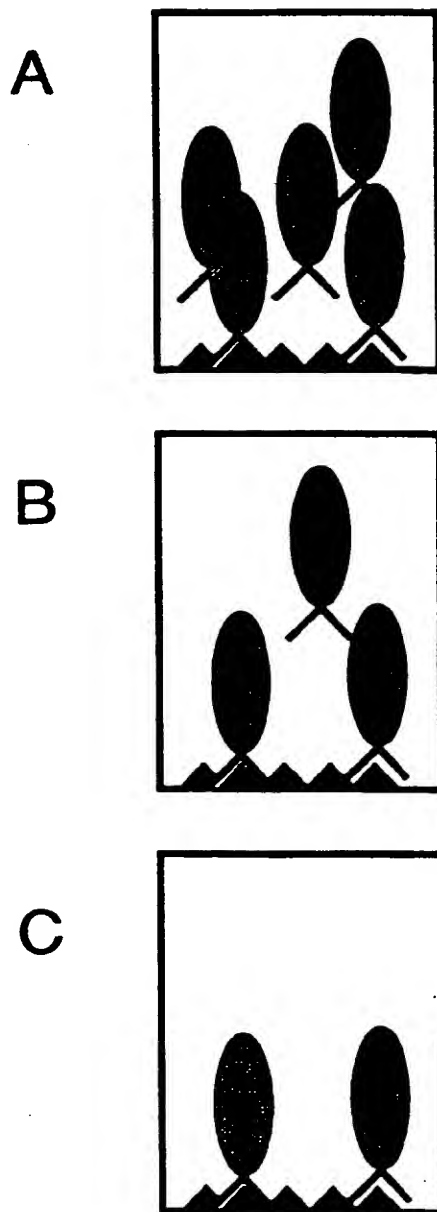


Fig. 13/14

```

1 EVQLVESGRGLVQPGGSLRLSCAASGFTFSHFLVAWVRQAPGKGLEWVAT 50
  ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1 EVQLVESGGGLVQPGGSLRLSCAASGFTFSXXXXXWVRQAPGKGLEWVAX 50

51 YSMISRARVLDGSFNTRYTISRDDSKNTLYLQMNSLRAEDTAVYYCARIG 100
  |:||||| ||||| ||||| ||||| ||||| ||||| |||||
51 XXXXXXXXXXXXXXXXXXXXRFTISRDDSKNTLYLQMNSLRAEDTAVYYCARXX 100

101 STHTIPRLSQYGGQGLVTVSSGGGGSGGGSGGGGSDIQMTQSPSSLSA 150
  ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
101 XXXXXXXXXXXXWGQGLVTVSSGGGGSGGGSGGGGSDIQMTQSPSSLSA 150

151 SVGDRVITCKLRGPQPHAITWYQQKPGKAPKLLIYYDQGLVGVPSRFS 200
  ||||| ||||| ||||| ||||| ||||| ||||| |||||
151 SVGDRVITCXXXXXXXXXXWYQQKPGKAPKLLIYXXXXXXGVPSRFS 200

201 GSGSGTDFTPTISSLEPEDFATYYCTPTHKIDSPFGQGTKVEIKRT 246
  ||||| ||||| ||||| ||||| ||||| ||||| |||||
201 GSGSGTDFTLTISLQPEDFATYYCXXXXXXXXXXFGQGTKVEIKRT 246

```

## Fig. 14/14

```

91 GAAGTTCAACTGGTTGAATCCGGTCGTGGTCTGGTTCAACCAGGTGGTTC 140
   |||||
1  gaagttcaactggttgaatccggtggtggtctggttcaaccaggtggttc 50

141 CCTGCGTCTGTCTCTGTGCTGCTTCCGGTTTCACCTTCTCCCATTTTTTGG 190
   |||||
51 cctgcgctctgtcctgtgctgcttccggtttcaccttctccnnnnnnnnnn 100

191 TGGCGTGGGTTCGTCAAGCTCCAGGTAAAGGTCTGGAATGGGTGCTACC 240
   ::::|
101 nnnnntgggttcgtcaagctccaggtaaaggtctggaatgggttgctnnn 150

241 TACTCAATGATTAGCCGGGCCCGaGTACTCGATGGCTCCTTTAATGGACG 290
   ::::|
151 nnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnncg 200

291 TTaACCATCTCCCGTGACGACTCCAAAAACACCCtgTACctgcaaataga 340
   || |||||
201 tttcaccatctcccgtgacgactccaaaaacaccctgtacctgcaaataga 250

341 actccctgcggtgctgaagaCACCGCTGtTTACTACTGTGCTCGTATTGGT 390
   |||||
251 actccctgcggtgctgaagacaccgctgtttactactgtgctcgtnnnnnn 300

391 TCTACGCACACAATCCCACGACTGTCTCAATACGGGGgTCAAGGTACCCT 440
   ::::|
301 nnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnntggggtcaaggtaccct 350

441 GGTtaccgtttcctccggtggtggtggttccggtggtgGTGGTTCTGGTG 490
   |||||
351 ggttaccgtttcctccggtggtggtggttccggtggtggtggttctggtg 400

491 GTGGTGGTTCCGACATCCAAATGACCCAATCCCCATCCTCTCTGTCCGCT 540
   |||||
401 gtggtggttccgacatccaaatgaccaatccccatcctctctgtccgct 450

541 TCCGTTGGTGACCGTGTTACCATCACCTGTAAACTCAGAGGACCACAACC 590
   |||||
451 tccgttgggtgaccgtgttaccatcacctgtnnnnnnnnnnnnnnnnnnnn 500

591 ACACGCCATTACATGGTACCAACAAAAACCAGGTAAAGCTCCAAAACCTGC 640
   ::::|
501 nnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnntggtaccaacaaaaaccaggtaaagctccaaaactgc 550

641 TGATctactACGACGGCCAAACGTTGGTGGGTGTTCCATCCCGTTTCTCC 690
   |||||
551 tgatctacnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnnggtgttccatcccggtttctcc 600

691 GGTTCTGGTCTCTGGTACCGACTTCACCCCGACCATCTCCTCTCTGGAACC 740
   |||||
601 ggttccggttctggtaccgacttcaccctgaccatctcctctctgcaacc 650

741 AGAAGACTTCGCTACCTACTACTGTACTCCTACGCACAAGATCGATAGCC 790
   |||||
651 agaagacttcgctacctactactgtnnnnnnnnnnnnnnnnnnnnnnnnnnnn 700

791 CATTCGGTCAAGGTACCAAAGTTGaAaTCAAACGTACC 828
   ::||
701 nnttcggtcaaggtaccaaagttgaaatcaaacgtacc 738

```



European Patent  
Office

## EUROPEAN SEARCH REPORT

Application Number

DOCUMENTS CONSIDERED TO BE RELEVANT			EP 93102609.0
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. CL.5)
A	<u>WO - A - 91/19 739</u> (CELLTECH LIMITED) * Claims * --	1, 9, 19, 20	C 12 P 21/00 C 07 K 13/00 C 12 N 15/09 C 12 N 15/70
A	<u>WO - A - 91/13 909</u> (PROTEUS MOLECULAR DESIGN LIMITED) * Claims 1, 27, 28, 32 * --	1, 12, 13, 25, 26	C 12 N 1/21 G 01 N 33/53 /(C 12 N 1/21 C 12 R 1:19)
A	<u>WO - A - 92/02 551</u> (B.R. CENTRE LIMITED et al.) * Claim 2 * ----	1	
			TECHNICAL FIELDS SEARCHED (Int. CL.5)
			C 12 P C 07 K C 12 N G 01 N
The present search report has been drawn up for all claims			
Place of search VIENNA		Date of completion of the search 07-06-199 .	Examiner WOLF
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons ----- Δ : member of the same patent family, corresponding document	

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